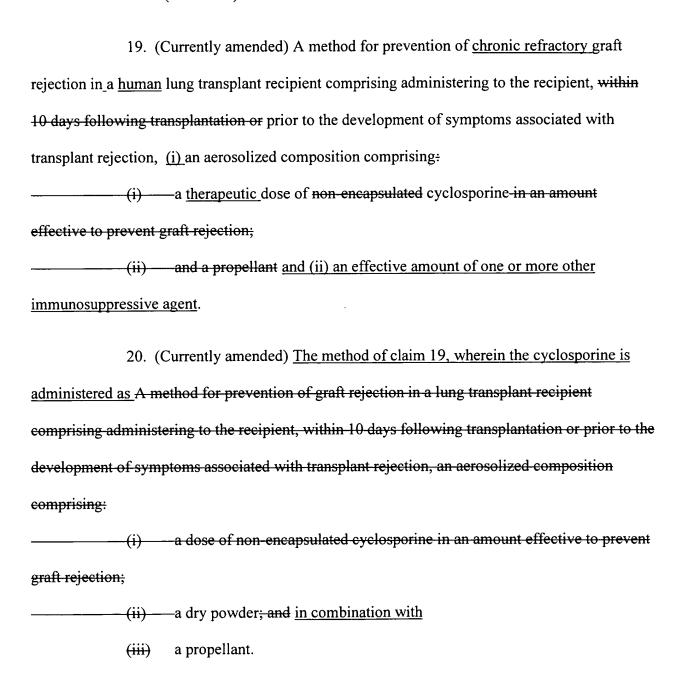
AMENDMENTS TO THE CLAIMS

The listing of claims provided below will replace all prior versions, and listings, of claims in the application.

Listing of Claims

1-18. (Cancelled)



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21. (Currently amended) The method of claim 19, wherein the cyclosporine is
dissolved in -A method for prevention of graft rejection in a lung transplant recipient comprising
administering to the recipient, within 10 days following transplantation or prior to the
development of symptoms associated with transplant rejection, an aerosolized composition
comprising:
(i) a dose of non-encapsulated cyclosporine in an amount
effective to prevent graft rejection;
————————an organic solvent ; and
——————————————————————————————————————
22. (Previously presented) The method of claim 19, 20 or 21 wherein the dose of
cyclosporine is sufficient to achieve deposition levels ranging between 15 and 30 mg in a lung.
23. (Cancelled)
24. (Currently amended) The method of claim 19, 20 or 21 wherein the
aerosolized composition is co-administered with [[a]] an anti-inflammatory reagent.
25. (Currently amended) A method for ameliorating pulmonary inflammation in a
subject having a lung disorder selected from the group consisting of asthma, sarcoidosis,
emphysema, cystic fibrosis, idiopathic pulmonary fibrosis, chronic bronchitis, hypersensitivity
pneumonitis and eosinophilic pneumonia, comprising administering to the subject an aerosolized
composition comprising-
(i) a dose of non-encapsulated cyclosporine dissolved in an organic solvent,
in an amount effective to inhibit or ameliorate pulmonary inflammation; and
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26. (Currently amended) A method for ameliorating pulmonary inflammation in a subject comprising administering to the subject an aerosolized composition comprising: a dose of non-encapsulated cyclosporine in dry powder form in an amount (i) effective to inhibit or ameliorate pulmonary inflammation; and a dry-powder; and (ii) a propellant. (iii) 27. (Cancelled) 28. (Cancelled) 29. (Currently amended) The method of claim 25[[,]] or 26 or 27 wherein the dose of cyclosporine is sufficient to achieve deposition levels ranging between 5 and 30 mg in a lung. 30. (Currently amended) A method for prevention of graft rejection in a non-lung transplant recipient comprising administering to the non-lung transplant recipient, within 10 days following transplantation or prior to the development of symptoms associated with transplant rejection, an aerosolized composition comprising: (i)—a dose of non-encapsulated cyclosporine dissolved in an organic solvent, in an amount effective to prevent graft rejection; (ii) and a propellant. 31. (Currently amended) A method for prevention of graft rejection in a nonlung transplant recipient comprising administering to the non-lung transplant recipient, within 10

days following transplantation or prior to the development of symptoms associated with

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transplant rejection, an aerosolized composition comprising:

- (i) a dose of non-encapsulated cyclosporine in dry powder form in an amount effective to prevent graft rejection; and
 - (ii) a dry powder; and
 - (iii) and a propellant.
 - 32. (Cancelled)
- 33. (Currently amended) A method for inhibiting the immune response associated with a T-cell mediated immune disorder in a subject, where said disorder is selected from the group consisting of type IV cell-mediated hypersensitivity, systemic lupus erythematosis, myasthenia gravis, Grave's disease, Hashimoto's thyroiditis, rheumatoid arthritis, scleroderma, and pernicious anemia, comprising administering, to the subject, to the non-lung transplant recipient, within 10 days following transplantation or prior to the development of symptoms associated with transplant rejection, an aerosolized composition comprising:

 (i)—a dose of non-encapsulated cyclosporine dissolved in an organic solvent, in an amount effective to inhibit the immune response associated with a T-cell mediated immune
- (ii) a propellant.

disorder; and

- 34. (Currently amended) The method of claim 30[[,]] or 31 or 32 wherein the dose of cyclosporine is sufficient to achieve circulating levels ranging between 50-250 ng/ml.
- 35. (Currently amended) The method of claim 30[[,]] or 31 or 32 wherein the aerosolized composition is co-administered with a second immunosuppressive agent.
- 36. (Currently amended) A method for inhibiting the immune response associated with a T-cell mediated immune disorder in a subject subject, where said disorder is selected from

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the group consisting of type IV cell-mediated hypersensitivity, systemic lupus erythematosis, myasthenia gravis, Grave's disease, Hashimoto's thyroiditis, rheumatoid arthritis, scleroderma, and pernicious anemia, comprising administering to the subject non-lung transplant recipient, within 10 days following transplantation or prior to the development of symptoms associated with transplant rejection, an aerosolized composition comprising:

(i) a dose of non-encapsulated cyclosporine <u>in dry powder form</u> in an amount effective to inhibit the immune response associated with a T-cell mediated immune disorder; <u>and</u>

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- (ii) a dry powder; and
- (iii) a propellant.

37-48. (Cancelled)

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